

**K972401 HOWMEDICA MODULAR REPLACEMENT SYSTEM -
DISTAL FEMORAL/PROXIMAL TIBIAL SEGMENTS**Jul 13, 1998
382 days to decisionK972401 · Product code: **KRO** · Orthopedic
Source: <https://www.510kdatabase.net/k972401/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Constrained, Cemented, Metal/polymer (KRO)
Date received	Jun 26, 1997
Decision date	Jul 13, 1998
Days to decision	382 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Howmedica Corp.
Location	Mchenry, IL, US
Contact	MARGARET F CROWE
510(k) history	373 submissions · 325 cleared · 1976-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k972401/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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